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Amendments to the Claims

Please amend the claims as follows:

Claim 1 (Currently Amended): A compound of the formula

$$(R^3)_2$$
 R^{11}
 R^2
 $(R^{3B})_2$
 $(R^{3B})_2$
 $(R^{3A})_2$
 $(R^{3A})_2$
 $(R^{3A})_2$
 $(R^{3A})_2$

or a pharmaceutically acceptable salt, solvate or ester thereof, wherein:

- (A) R¹ is selected from the group consisting of:
 - (1) unsubstituted aryl; and
 - (2) anyl substituted with one or more R⁵ groups;
- (3) unsubstituted heteroaryl; and
- ———(4) heteroaryl-substituted with one-or-more R⁵ groups,
- (B) R² is selected from the group consisting of:
 - (1) alkyl;
 - (2) -XC(O)Y;
 - (3) $-(C_1-C_6)$ alkylene-XC(O)Y;
 - (4) $-(C_0-C_6)$ alkylene- (C_3-C_6) cycloalkylene- (C_0-C_6) alkylene-XC(O)Y;
 - (5) aryl;
 - (6) aryl substituted with one or more R⁵ groups;
 - (7) heteroaryl:
 - (8) heteroaryl substituted with one or more R⁵ groups;
 - (91) cycloalkylene-X-C(O)-Y;
 - (492) -CH₂-X-C(O)-NR³-Y;
 - (443) -CH2-X-C(O)-Y; and
 - (424) -CH2-X-C(O)-NR3-Y,
- (C) Each R³ is independently selected from the group consisting of:
 - (1) H; and
 - (2) alkyl,
- (D) Each R^{3A} and R^{3B} is independently selected from the group consisting of:

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- (1) H; and
- (2) alkyl;
- (E) R⁵ is independently selected from the group consisting of:
 - (1) halo;
 - (2) -CF₃;
 - (3) -OH;
 - (4) -O-alkyl;
 - (5) -OCF₃;
 - (6) -CN;
 - (7) -NH₂;
 - (8) -C(O)₂alkyl;
 - (9) -C(O)NR⁶R⁷;
 - (10) -alkylene-NR⁶R⁷;
 - (11) -NR⁶C(O)alkyl;
 - (12) -NR⁶C(O)aryl;
 - (13) -NR⁶C(O)heteroaryl; and
 - (14) $-NR^6C(O)NR^6R^7$;
- (F) X is selected from the group consisting of:
 - (1) -O-;
 - (2) -NH-;
 - (3) -N-alkyl; and
 - (4) -O-alkylene;
- (G) Y is selected from the group consisting of:
 - (1) $-NR^6R^7$;
 - (2) $-N(R^3)(CH_2)_bNR^6R^7$ wherein b is 2-6;
 - (3) unsubstituted aryl;
 - (4) unsubstituted heteroaryl;
 - (5) -alkyl;
 - (6) -cycloalkyl,
 - (7) unsubstituted arylalkyl;
 - (8) unsubstituted arylcycloalkyl;
 - (9) unsubstituted heteroarylalkyl;
 - (10) unsubstituted heteroarylcycloalkyl;
 - (11) unsubstituted arylheterocycloalkyl;

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- (12) substituted aryl;
- (13) substituted heteroaryl;
- (14) substituted arylalkyl;
- (15) substituted arylcycloalkyl;
- (16) substituted heteroarylalkyl;
- (17) substituted heteroarylcycloalkyl; and
- (18) substituted arylheterocycloalkyl;
- (19) substituted heterocycloalkyl alkyl;
- (20) unsubstituted heteroaryl alkyl;
- (21) unsubstituted aryl alkyl heterocycloalkyl;
- (22) unsubstituted heterocycloalkyl; and
- (23) unsubstituted cycloalkyl,

wherein the aryl moiety in said substituted groups (12), (14), (15), (18), and (21) of said Y group, and the heteroaryl moiety in said substituted groups (13), (16), (17) and (20) of said Y group, are substituted with one or more substituents independently selected from the group consisting of:

- (a) halo;
- (b) -CF₃;
- (c) -OH;
- (d) -O-alkyl;
- (e) -OCF₃;
- (f) -CN;
- (g) -NH₂;
- (h) $-C(O)_2(C_1-C_6)$ alkyl;
- (i) $-C(O)NR^6R^7$;
- (j) $-(C_1-C_6)$ alkylene-NR⁶R⁷;
- (k) -NR⁶C(O)alkyl;
- (I) -NR⁶C(O)aryl;
- (m) -NR⁶C(O)heteroaryl;
- (n) -NR⁶C(O)NR⁶R⁷; and
- (o) alkyl,

or Y is selected from the group consisting of:

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$$(C) \qquad (R^8)_r \qquad (R^8)_r$$

- (H) R⁶ and R⁷ are independently selected from the group consisting of:
 - (1) H;
 - (2) alkyl;
 - (3) cycloalkyl;
 - (4) arylalkyl;
 - (5) heteroarylalkyl;
 - (6)

(7)

$$\begin{cases} (R^8)_s \\ N-R^9 \end{cases}$$
 (b) ; and

- (8) heterocycloalkyl,
- (I) Each R⁸ is independently selected from the group consisting of:
 - (1) alkyl;
 - (2) alkyl substituted with 1 to 4 hydroxy groups; and
 - (3) -OH,

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- (J) Each R⁹ is independently selected from the group consisting of:
 - (1) H;
 - (2) alkyl;
 - (3) alkyl substituted with 1 to 4 hydroxy groups;
 - (4) cycloalkyl;
 - (5) cycloalkyl substituted with 1 to 4 hydroxy groups;
 - (6) arylalkyl;
 - (7) heteroarylalkyl;
 - (8) -C(O)O-alkyl;
 - (9) alkylene-O-alkylene-OH;
 - (10) aryl substituted with one or more R⁵ groups;
 - (11) heteroaryl substituted with one or more R⁵ groups;
 - (12) unsubstituted heteroaryl;
 - (13) unsubstituted aryl;
 - (14) -alkylene-C(O)O-alkyl; and
 - (15) hydroxyalkyl-O-alkyl,
- (K) Each R¹⁰ is independently selected from the group consisting of:
 - (1) H; and
 - (2) alkyl,
- (L) R¹¹ is selected from the group consisting of:
 - (1) unsubstituted aryl;
 - (2) substituted aryl;
 - (3) unsubstituted heteroaryl,
 - (4) alkyl;
 - (5) cycloalkyl;
 - (6) unsubstituted arylalkyl;
 - (7) unsubstituted arylcycloalkyl,
 - (8) unsubstituted heteroarylalkyl;
 - (9) unsubstituted heteroarylcycloalkyl;
 - (10) unsubstituted arylheterocycloalkyl;

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- (11) alkoxyalkyl;
- (12) substituted heteroaryl;
- (13) substituted arylalkyl;
- (14) substituted arylcycloalkyl;
- (15) substituted heteroarylalkyl; and
- (16) substituted arylheterocycloalkyl,

wherein the aryl moiety in said substituted groups (2), (13), (14) and (16) of said R¹¹ group, and the heteroaryl moiety in said substituted groups (12) and (15) of said R¹¹ group, are substituted with one or more substituents independently selected from the group consisting of:

- (a) halo;
- (b) -CF₃;
- (c) -OH;
- (d) -O-alkyl;
- (e) -OCF₃;
- (f) -CN;
- (g) -NH₂;
- (h) $-C(O)_2(C_1-C_6)$ alkyl;
- (i) $-C(O)NR^6R^7$;
- (j) $-(C_1-C_0)$ alkylene-NR⁶R⁷;
- (k) -NR⁶C(O)alkyl;
- (I) -NR⁶C(O)aryl;
- (m) -NR⁶C(O)heteroaryl; and
- (n) $-NR^6C(O)NR^6R^7$;
- (M) (1) m is 0 to 3, and if m is greater than 1, m moieties can be the same or different from one another;
- (2) n is 0 to 3, and if n is greater than 1, n moieties can be the same or different from one another;
- (3) o is 0 to 3, and if o is greater than 1, o moieties can be the same or different from one another;
- such that m+n+o is 1, 2, 3 or 4,
- (N) p is 0 to 4, and if greater than 1, p moieties can be the same or different from one another;

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- (O) r is 0 to 4, and if greater than 1, r moieties can be the same or different from one another;
- (P) s is 0 to 3, and if greater than 1, s moleties can be the same or different from one another; and
- (Q) Z is selected from the group consisting of:
 - unsubstituted heterocycloalkyl;
 - (2) substituted heterocycloalkyl;
 - (3) -NH₂;
 - (4) -NH(alkyl);
 - (5) -N(alkyl)₂ wherein each alkyl is the same or different;
 - (6) -NH(unsubstituted cycloalkyl);
 - (7) -NH(substituted cycloalkyl);
 - (8) -N(alkyl)(unsubstituted cycloalkyl);
 - (9) -N(alkyl)(substituted cycloalkyl);
 - (10) -NH(unsubstituted aralkyl);
 - (11) -NH(substituted aralkyl);
 - (12) -N(alkyl)(aralkyl);
 - (13) -NH(unsubstituted heterocycloalkyl);
 - (14) -NH(substituted heterocycloalkyl);
 - (15) -N(alkyl)(unsubstituted heterocycloalkyl),
 - (16) -N(alkyl)(substituted heterocycloalkyl);
 - (17) -NH(unsubstituted heteroaralkyl);
 - (18) -NH(substituted heteroaralkyl);
 - (19) -NH-alkylene-(unsubstituted cycloalkyl);
 - (20) -NH-alkylene-(substituted cycloalkyl);
 - (21) -N(alkyl)alkylene-(unsubstituted cycloalkyl);
 - (22) -N(alkyl)alkylene-(substituted cycloalkyl);
 - (23) -NHalkylene-(unsubstituted heterocycloalkyl);
 - (24) -NHalkylene-(substituted heterocycloalkyl);
 - (25) -N(alkyl)alkylene-(unsubstituted heterocycloalkyl);
 - (26) -N(alkyl)alkylene-(substituted heterocycloalkyl);
 - (27) unsubstituted benzofused heterocycloalkyl; and
 - (28) substituted benzofused heterocycloalkyl;
 - (29) H; and

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(30) -N(hydroxyalkyl)₂, wherein each alkyl may be the same or different,

wherein said substituted heterocycloalkyl moiety of substituents (2), (14), (16), (24), (26) and (27) of group Z, and said substituted cycloalkyl moiety of substituents (7), (9), (20) and (22) of group Z, and said substituted aryl moiety of substituent (11) of group Z, and said substituted heteroaryl moiety of substituent (18) of group Z, are substituted with 1 to 3 groups independently selected from the group consisting of:

- (a) alkyl;
- (b) -OH;
- (c) -Oalkyl;
- (d) -OC(O)alkyl;
- (e) -OC(O)aryl;
- (f) -NH₂;
- (g) -NH(alkyl);
- (h) -N(alkyl)₂ wherein each alkyl is the same or different;
- (i) -NHC(O)alkyl;
- (j) -N(alkyl)C(O)alkyl;
- (k) -NHC(O)aryl;
- (I) -N(alkyl)C(O)aryl;
- (m) -C(O)alkyl;
- (n) -C(O)aryl;
- (o) -C(O)NH₂;
- (p) -C(O)NH(alkyl);
- (q) -C(O)N(alkyl)₂ wherein each alkyl is the same or different;
- (r) -C(O)2alkyi;
- (s) -alkylene-C(O)Oalkyl;
- (t) piperidinyl;
- (u) pyrrolidinyl;
- (v) 1,1-ethylenedioxy;
- (w) aryl;
- (x) heteroaryl; and
- (y) -O-CH₂CH₂-O-wherein both oxygen atoms are bound to the same carbon atom, and provided that the aryl and heteroaryl

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moieties of said Z group are not substituted with said -O-CH₂CH₂-O- group.

Claim 2 (Currently Amended): The compound of Claim 1 wherein:

- (A) R¹ is anyl substituted with one or more R⁵ groups;
- (B) n is 0 or 1 and m is 1, 2 or 3 such that m+n is 3;
- (C) p is 0 or 1; and
- (D) R^2 is -XC(O)Y , -(C₁-C₆)alkylene-XC(O)Y or -(C₀-C₆)alkylene-(C₃-C₆)cycloalkylene-(C₀-C₆)alkylene-XC(O)Y.

Claim 3 (Original): The compound of Claim 2 wherein:

- (A) R¹ is phenyl substituted with one or more R⁵ groups; and
- (B) n is 0 and m is 3.

Claim 4 (Currently Amended): The compound of Claim 1, wherein R^2 is X-C(O)-Y

Claim 5 (Original): The compound of Claim 3 wherein R¹ is phenyl substituted with one or more halo atoms.

Claim 6 (Currently Amended): The compound of Claim 1 wherein:

- (A) R¹ is anyl substituted with one or more R⁵ groups;
- (B) n is 0 or 1 and m is 1, 2 or 3 such that m+n is 3;
- (C) p is 0 or 1;
- (D) R^2 is -XC(O)Y, -(C₁-C₆)alkylene-XC(O)Y or -(C₀-C₆)alkylene-(C₃-C₆)cycloalkylene-(C₀-C₆)alkylene-XC(O)Y;
- (E) X is O;
- (F) Y is -NR⁶R⁷; or Y is selected from the group consisting of:

$$(c) \qquad (d) \qquad (e) \qquad (f)$$

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$$(R^{10})_p$$
 and
$$(R^{10})_p$$
 and
$$(R^{10})_p$$
 ; and

(G) R^6 and R^7 are independently selected from the group consisting of: H, methyl, ethyl, -(C₃-C₆)cycloalkyl, -aryl(C₁-C₆)alkyl, 4-pyridylmethyl, and

$$\begin{cases} (R^8)_r & \text{and} & \begin{cases} (R^8)_s \\ N - R^9 \end{cases} \end{cases}$$

Claim 7 (Original): The compound of Claim 6 wherein:

- (A) R¹ is phenyl substituted with one or more R⁵ groups;
- (B) n is 0 and m is 3;
- (C) said group

is a group of the formula:

(D) said group

is a group of the formula:

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(E) R^{11} is selected from the group consisting of: -(C₁-C₆)alkyl, (C₃-C₈)-cycloalkyl, aryl, aryl(C₁-C₆)alkyl and -(C₁-C₆)alkoxyalkyl.

Claim 8 (Original): The compound of Claim 7 wherein said R¹¹ is selected from the group consisting of: methyl, ethyl, cyclohexyl, phenyl, benzyl, –(CH₂)₂phenyl, and –CH₂OCH₃.

Claim 9 (Original): The compound of Claim 7 wherein R¹ is phenyl substituted with one or more halo atoms.

Claim 10 (Original): The compound of Claim 8 wherein R¹¹ is phenyl substituted with one or more halo atoms.

Claim 11 (Original): The compound of Claim 6 wherein Y is selected from the group consisting of:

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Claims 12-14 (Cancelled).

Claim 15 (Original): A pharmaceutical composition comprising at least one compound of Claim 1 and at least one pharmaceutically acceptable carrier.

Claim 16 (Original): A method of inhibiting gamma-secretase in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one or more compounds of Claim 1.

Claim 17 (Original): A method of treating one or more neurodegenerative diseases in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one or more compounds of Claim 1.

Claim 18 (Original). A method of inhibiting the deposition of beta amyloid protein in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one more compounds of Claim 1.

Claim 19 (Original): A method of treating Alzheimer's disease in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one or more compounds of Claim 1.

Claim 20 (Original): A compound selected from the group consisting of:

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Claim 21 (Original): A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 20, or a pharmaceutically acceptable salt, ester or solvate thereof, together with a pharmaceutically acceptable excipient, diluent or carrier.

Claim 22 (Original): A method of inhibiting gamma-secretase in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one or more compounds of Claim 20.

Claim 23 (Original): A method of treating one or more neurodegenerative diseases in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one or more compounds of Claim 20.

Claim 24 (Original): A method of inhibiting the deposition of beta amyloid protein in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one more compounds of Claim 20.

Claim 25 (Original): A method of treating Alzheimer's disease in a patient in need of such treatment comprising administering to said patient a therapeutically effective amount of one or more compounds of Claim 20.

Claim 26 (Original): A compound of the following formula

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or a pharmaceutically acceptable salt, ester or solvate of said compound.

Claim 27 (Original): A compound of the following formula

or a pharmaceutically acceptable salt, ester or solvate of said compound.

Claim 28 (Original): A compound of the following formula

or a pharmaceutically acceptable salt, ester or solvate of said compound.

Claim 29 (Original): A compound of the following formula

or a pharmaceutically acceptable salt, ester or solvate of said compound.

Claim 30 (Original): A compound of the following formula

or a pharmaceutically acceptable salt, ester or solvate of said compound. 55775-1

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Claim 31 (Original): A compound of the following formula

or a pharmaceutically acceptable salt, ester or solvate of said compound.